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<!--StartFragment-->RESULT 2
AAD56890
ID   AAD56890 standard; cDNA; 1279 BP.
XX
AC   AAD56890;
XX
DT   06-NOV-2003  (first entry)
XX
DE   Human diacylglycerol acyltransferase 2 (DGAT2) cDNA, 112023.
XX
KW   Human; diacylglycerol acyltransferase 2; DGAT2; obesity; arrhythmia;
KW   coronary artery disease; hypertension; heart failure; tissue typing;
KW   aberrant lipogenesis; cardiovascular disorder; atherosclerosis; angina;
KW   atrial fibrillation; dilated cardiomyopathy; idiopathic cardiomyopathy;
KW   diabetes; chromosome mapping; forensic biology; enzyme; gene; ss.
XX
OS   Homo sapiens.
XX
FH   Key             Location/Qualifiers
FT   CDS             42. .1028
FT                   /*tag=  a
FT                   /product= "Human diacylglycerol acyltransferase 2"
XX
PN   WO2003053363-A2.
XX
PD   03-JUL-2003.
XX
PF   19-DEC-2002; 2002WO-US040974.
XX
PR   19-DEC-2001; 2001US-0341947P.
PR   19-SEP-2002; 2002US-0411859P.
XX
PA   (MILL-) MILLENNIUM PHARM INC.
XX
PI   Gimeno RE,  Wu Z,  Kapeller-Libermann R,  Hubbard BK;
XX
DR   WPI; 2003-559092/52.
DR   P-PSDB; AAE37790.
XX
PT   New human diacylglycerol acyltransferase 2 (DGAT2) family member
PT   polypeptide and nucleic acid molecules, useful for diagnosing and
PT   treating obesity, diabetes, atherosclerosis, aberrant lipogenesis or
PT   triglyceride synthesis.
XX
PS   Claim 1; Page 133-134; 154pp; English.
XX
CC   The invention relates to human diacylglycerol acyltransferase 2 (DGAT2)
CC   family members and their uses. DGAT2 family member sequences or their
CC   modulators are useful for diagnosing and treating a subject with a
CC   disorder associated with the aberrant DGAT family member polypeptide
CC   activity or nucleic acid expression, such as a disorder associated with
CC   obesity, diabetes, aberrant lipogenesis or triglyceride synthesis, or
CC   cardiovascular disorder (e.g. atherosclerosis, coronary artery disease,
CC   hypertension, heart failure, atrial fibrillation, arrhythmias, dilated
CC   cardiomyopathy, idiopathic cardiomyopathy or angina). The invention is
CC   also useful in screening assays (e.g. tissue typing, chromosome mapping,
CC   or in forensic biology), in predictive medicine (e.g. diagnostic assays,
CC   prognostic assays, monitoring clinical trials or pharmacogenetics), or as
CC   surrogate markers (e.g. markers of disease states or markers of drug
CC   activity). The present sequence is human DGAT2 cDNA
XX
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SQ   Sequence 1279 BP; 273 A; 352 C; 328 G; 326 T; 0 U; 0 Other;

  Query Match             95.9%;  Score 1084.8;  DB 2;  Length 1279;
  Best Local Similarity   98.0%;
  Matches 1109;  Conservative    0;  Mismatches   22;  Indels    1;  Gaps    1;

Qy          1 ACTGTTCTGAGATCTTTGCCTCCCTCAGGCTCCCGAGAATCATGGCTCATTCCAAGCAGC 60
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Qy         61 CTAGTCACTTCCAGAGTCTGATGCTTCTGCAGTGGCCTTTGAGCTACCTTGCCATCTTTT 120
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Qy        181 TTTACTTTGCCTGGTTGTTCCTGGACTGGAAGACCCCAGAGCGAGGTGGCAGGCGTTCGG 240
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Qy        601 AAAGTGTGCCCAACACCACCACCCTCATCCTCCAGAAGCGCAAGGGGTTCGTGCGCACAG 660
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Qy        661 CCCTCCAGCATGGGGCTCATCTGGTCCCCACCTTCACTTTTGGGGAAACTGAGGTGTATG 720
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Qy        721 ATCAGGTGCTGTTCCATAAGGATAGCAGGATGTACAAGTTCCAGAGCTGCTTCCGCCGTA 780
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Db        721 ATCAGGTGCTGTTCCATAAGGATAGCAGGATGTACAAGTTCCAGAGCTGCTTCCGCCGTA 780

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